MEDISTIK ICE- menthol, camphor solution
Natureteq Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

NATURETEQ - MEDISTIK PRO ICE THERAPY, 50231-411

ACTIVE INGREDIENTS

Menthol 10%

Camphor 4%

PURPOSE

Topical Analgesic

USES

For the temporary relief of sore muscles and joints associated with

- strains and sprains
- backaches
- lumbago
- pain of tendons and/or ligaments
- arthritic or rheumatic pain

WARNINGS

For external use only. Avoid contact with eyes and mucous membranes. Do not apply to wounds or damaged skin. Do not tightly bandage. Do not apply with external heat, such as an electric heating pad, as this may result in excessive skin irritation or skin burn.

Stop use and consult a physician if condition worsens, rash or irritation develops, or if symptoms persist for more than 7 days or clear up and recur in a few days. Consult a health care practitioner prior to use if you are pregnant or breastfeeding. Rashes and/or burning discomfort, and hypersensitivity such as allergy have been known to occur; in which case, discontinue use.

DIRECTIONS

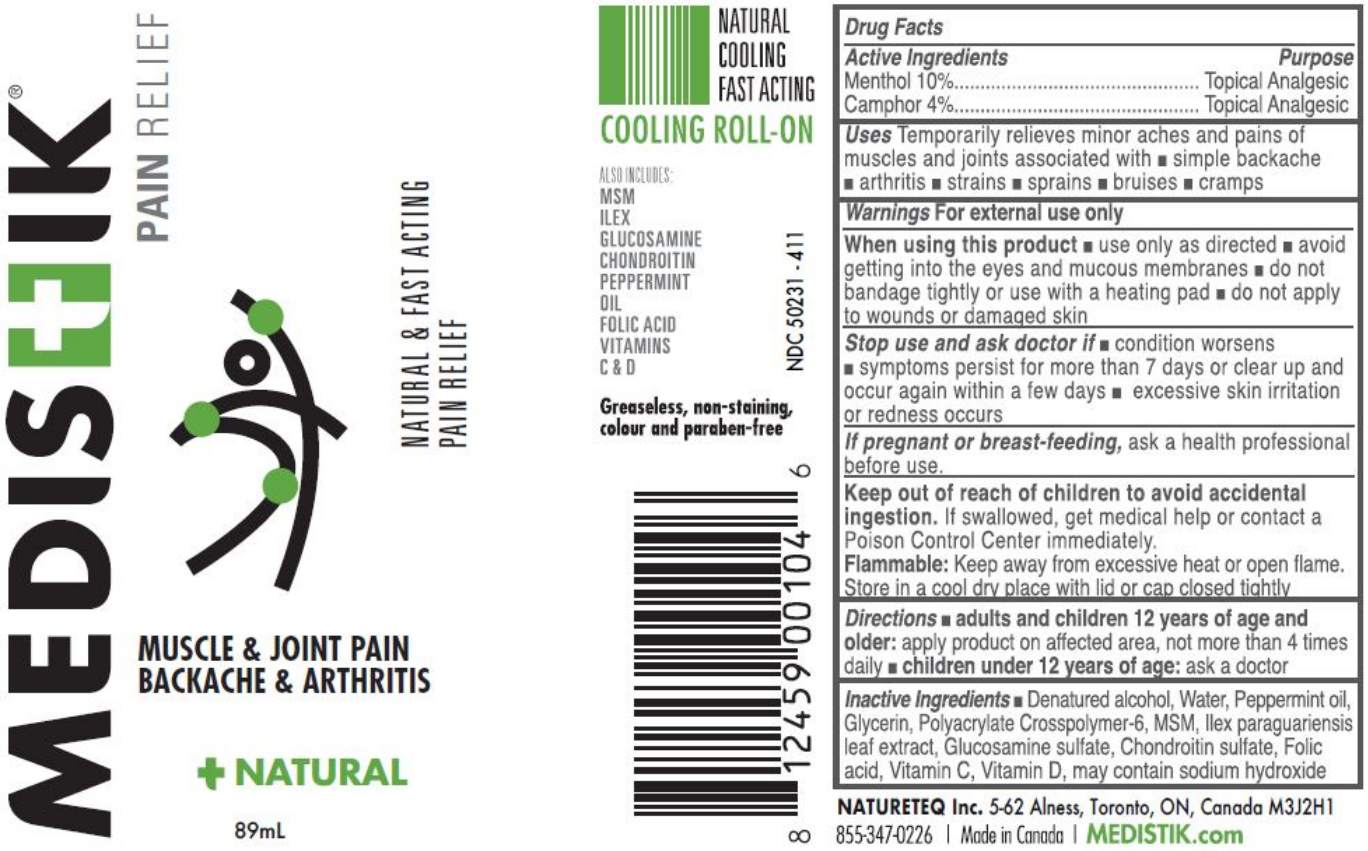
For use on adults and children over 12 years of age.

Apply to affected area(s) 3 to 4 times per day as required. A stinging or burning sensation will be experienced during the first few minutes as the formula begins working. For arthritis or muscle pain of the hands, retain for at least 10 minutes then wash hands.

INACTIVE INGREDIENTS

Denatured alcohol, Water, Peppermint oil, Glycerin, Acrylates copolymer, MSM, Ilex paraguariensis leaf extract, Glucosamine sulphate, Chondroitin sulphate, Folic acid, Vitamin C, Vitamin D.

Cautions: Keep out of reach of children. If overdose or accidental ingestion occurs, call a Poison Control Center immediately.



MEDISTIK ICE

menthol, camphor solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50231-411
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10 g in 100 mL
CAMPHOR OIL (UNII: 75IZZ8Y727) (CAMPHOR OIL - UNII:75IZZ8Y727)	CAMPHOR OIL	4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
CHONDROITIN SULFATE (CHICKEN) (UNII: 7VZ9466BAB)	
FOLIC ACID (UNII: 935E97BOY8)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
VITAMIN D (UNII: 9VU1KI44GP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50231-411-11	89 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/21/2020	
2	NDC:50231-411-12	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/18/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	02/17/2016	

Labeler - Natureteq Inc. (243737371)

Revised: 11/2021 Natureteq Inc.